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PATENT

- 24. A pharmaceutical composition comprising the polypeptide of claim 21 and a pharmaceutically acceptable carrier.
- 25. An immunogenic composition comprising the polypeptide of claim 21 and an adjuvant.
- The composition of claim 25, wherein the adjuvant is selected from the group consisting of Freund's adjuvant, a mineral gel, aluminum hydroxide, lysolecithin, pluronic polyol, a polyanion, a peptide, an oil emulsion, keyhole limpet hemocyanin (KLH), dinitrophenol (DNI), Bacillus Calmette-Guerin, and Corynebacterium parvum.
- 27. A method for eliciting an immune response to telomerase reverse transcriptase protein in a subject, comprising administering to the subject the polypeptide of claim 21.
- 28. The method of claim 27, further comprising assessing whether telomerase-specific antibody is produced as a result of the administration.
- 29. An immunogenic composition comprising a polypeptide consisting essentially of 5 to 10 consecutive amino acids of the sequence provided in SEQ. ID NO:225, and an adjuvant.
- 30. The composition of claim 29, wherein the adjuvant is selected from the group consisting of Freund's adjuvant, a mineral gel, aluminum hydroxide, lysolecithin, pluronic polol, a polyanion, a peptide, an oil emulsion, keyhole limpet hemocyanin (KLH), dinitrophenol (DNF), Bacillus Calmette-Guerin, and Corynebacterium parvum.
- 31. A method for eliciting an immune response to telomerase reverse transcriptase protein in a subject, comprising administering to the subject a polypeptide consisting essentially of 5 to 10 consecutive amino acids of the sequence provided in SEQ. ID NO:225.
- 32. The method of claim 31, further comprising assessing whether telomerase-specific antibody is produced as a result of the administration.

